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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/692,083	10/22/2003	William Martin Belef	704117.4005	8282	
ORRICK, HER IP PROSECUT	7590 05/25/200 RINGTON & SUTCL ION DEPARTMENT			EXAMINER GANESAN, SUBA	
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)				
	10/692,083 ·	BELEF ET AL.				
Office Action Summary	Examiner	Art Unit				
	Suba Ganesan	3738				
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address				
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tim rill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).				
Status						
1)⊠ Responsive to communication(s) filed on 01 M	arch 2007.	•				
3) Since this application is in condition for allowar	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is					
closed in accordance with the practice under E	x parte Quayle, 1935 C.D. 11, 45	53 O.G. 213.				
Disposition of Claims						
4) Claim(s) 1-33 is/are pending in the application.						
4a) Of the above claim(s) is/are withdrawn from consideration.						
	5) Claim(s) is/are allowed.					
6)⊠ Claim(s) <u>1-33</u> is/are rejectéd. 7)□ Claim(s) is/are objected to.						
8) Claim(s) are subject to restriction and/or	r election requirement.					
	•					
Application Papers						
9) The specification is objected to by the Examiner						
10) The drawing(s) filed on is/are: a) accepted or b) objected to by the Examiner. Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).						
Replacement drawing sheet(s) including the correcti	·					
11) The oath or declaration is objected to by the Ex	· · · · · · · · · · · · · · · · · · ·	•				
Priority under 35 U.S.C. § 119						
12) Acknowledgment is made of a claim for foreign	priority under 35 U.S.C. § 119(a)	H-(d) or (f).				
a) ☐ All b) ☐ Some * c) ☐ None of:						
1. Certified copies of the priority documents have been received.						
2. Certified copies of the priority documents have been received in Application No						
_ ,	3. Copies of the certified copies of the priority documents have been received in this National Stage					
application from the International Bureau (PCT Rule 17.2(a)).						
* See the attached detailed Office action for a list of the certified copies not received.						
Attachment(s)						
1) D Notice of References Cited (PTO-892)	4) Interview Summary	(PTO-413)				
2) DNotice of Draftsperson's Patent Drawing Review (PTO-948)	Paper No(s)/Mail Da	nte				
Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date 5) Notice of Informal Patent Application 6) Other:						

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DETAILED ACTION

Response to Arguments

1. Applicant's arguments filed 3/1/2007 have been fully considered but they are not persuasive. Lambrecht discloses direct application of thermal energy to tissue (col. 29 lines 26-29, note that the RF energy directly heats surrounding tissue), resulting in tissue shrinkage and/or shrinkage of the defect (col. 20 lines 38-41).

Claim Rejections - 35 USC § 102

- 2. The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.
- 3. Claims **1-12** are rejected under 35 U.S.C. 102(b) as being anticipated by Lambrecht et al. (U.S. Pat. No. 6,482,235).

Lambrecht et al. discloses a method for augmenting an intervertebral disc in order to repair defects in the annulus fibrosis including creating an opening through the annulus fibrosis into the interior of the disc (see figure 19). Regarding claims 1-4, Lambrecht discloses removing a portion of the nucleus pulposus (col. 17 lines 4-7), introducing an implanted barrier (12). The barrier (12) is considered to be a therapeutic agent.

Lambrecht further discloses using radio frequency energy (col. 20, line 29) by introducing an 'elongate member' (130) with electrodes disposed on its distal portion, those electrodes being activated while the elongate member is within the interior of the

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disc. With respect to claim 6, a distal end of a needle is used to deliver barrier (12) (see fig. 29 A-D). A thermal device (30) is attached to the elongate member (130) that delivers electrical energy to the surrounding tissue in order to close the passage (see fig. 29 D). With respect to claim 12, Lambrecht discloses a secondary object (418) that is a handle member that has an electrically conductive filament (col. 27 lines 38-44).

Claim Rejections - 35 USC § 103

- 4. The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.
- 5. Claims **13** and **14** are rejected under 35 U.S.C. 103(a) as being unpatentable over Lambrecht '235 in view of Underwood et al. (U.S. Pat. No. 6,929,640).

Lambrecht is explained as above. However, Lambrecht does not disclose the injection of a therapeutic agent using a syringe, or disconnecting the syringe before connecting a handle member to the needle. Underwood discloses a device for closing openings in the interior of spinal discs using radio frequency energy transmitted via electrodes on the distal portion of an 'elongate element' (306). Underwood further discloses the use of a needle and syringe for the purpose of delivering saline solution to the area (see fig 16). Therefore it would have been obvious to one of ordinary skill in the art to modify Lambrecht with the syringe delivery system of Underwood for the purpose of delivering a therapeutic agent, such as an irrigant.

6. Claims **15-18**, **20,23**, and **27-29** are rejected under 35 U.S.C. 103(a) as being unpatentable over Froning (U.S. Pat. No.3,875,595) in view of Lambrecht '235.

Froning discloses a method of treating a spine, including the steps of removing at least portion of the nucleus pulposus from an interior region of a spinal disc to define a space; lining the space with a nonporous liner material or bladder (46); and filling the space with a fill material or fluid to expand the liner material (see Figures 1-8). The bladder includes a neck with an opening and a sealing member. However, Froning does not disclose the liner material being bioabsorbable, the use of energy to close the opening in the annulus fibrosis or the fill material used in the method being the nucleus pulposus from the disc.

Lambrecht teaches the use of resorbable materials such as polylactic acid and polyglycolic acid (for example see col. 11 lines 38-41). Lambrecht further teaches the use of radio frequency energy (col. 20 line 29) in order to close an opening in the annulus fibrosis. Therefore it would have been obvious to one of ordinary skill in the art to modify Froning with the radio frequency energy and bioabsorbable materials of Lambrecht in order to close an opening in the annulus fibrosis and to allow the lining to resorb over time.

With respect to claim 16, Lambrecht teaches the use of the nucleus pulposus within a defect (col. 21 line 38). It would have been obvious to one having ordinary skill in the art at the time the invention was made to practice the method of Froning with the fill material or fluid being the nucleus pulposus from the disc, since it has been held to be within the general skill of a worker in the art to select a known material on the basis of its suitability for the intended use as a matter of obvious design choice. In re Leshin, 125 USPQ 416. Moreover, applicant has not disclosed that use of the nucleus pulposus

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solves any stated problem or is for any particular purpose and it appears that the invention would perform equally well if the fill material did not have any nucleus pulposus from the disc.

Regarding claim 17, it would have been obvious to one of ordinary skill in the art to use the nucleus pulposus from the same patient in order to avoid homologous reactions. With respect to claim 18 and 28, it is known that using the nucleus pulposus means that naturally occurring extra-cellular matrix material is used (the natural material surrounding the chondrocyte-like cells of the nucleus pulposus). Furthermore, with respect to claim 20, nucleus pulposus from the same patient comprises an autologous therapeutic agent.

With regard to the fill material further having at least one of, a concentrated growth factor derived from centrifuged plasma of the patient (claim 21), it is noted that it would have been further obvious to one having ordinary skill in the art at the time the invention was made to practice the modified method of Froning with the fill material further having a concentrated growth factor derived from centrifuged plasma of the patient since it has been held to be within the general skill of a worker in the art to select a known material on the basis of its suitability for the intended use as a matter of obvious design choice. In re Leshin, 125 USPQ 416.

7. Claim 19, and 24-26 are rejected under 35 U.S.C. 103(a) as being unpatentable over Froning '595 in view of Lambrecht '235 as applied to claim 15 above, and further in view of Carr Jr. et al. (U.S. Pat. No. 5,733,337). Froning in view of Lambrecht discloses same as above. However, Froning in view of Lambrecht does not disclose a fill material

comprised of at least one of intestinal submucosa, stomach submucosa, or bladder submucosa. Carr Jr. et al. discloses the use of intestinal submucosa (col. 2-3, lines 66-3) for biodegradable implantation within the body. Therefore it would have been obvious to one of ordinary skill in the art to modify the method of Froning in view of Lambrecht to further include the use of intestinal submucosa for the purpose of making the implant biodegradable. It is also noted that intestinal submucosa comprises an extra-cellular matrix material.

- 8. Claim 22 is rejected under 35 U.S.C. 103(a) as being unpatentable over Froning '595 in view of Lambrecht '235 as applied to claim 15 above, and further in view of Felt et al. (U.S. Pat. No. 6,140,452). Froning in view of Lambrecht discloses same as above. However, Froning in view of Lambrecht does not disclose filler material comprising an interpenetrating polymer network material. Felt et al. teaches the use of an interpenetrating polymer network (for example, see col. 30 lines 36-40) in order to utilize a multiphasic bulk morphology. Therefore it would have been obvious to one of ordinary skill in the art to modify Froning in view of Lambrecht with the interpenetrating network of Felt et al. in order to have a filler material with multiphasic bulk morphology.
- 9. Claims **30-33** are rejected under 35 U.S.C. 103(a) as being unpatentable over Froning in view of Lambrecht '235 as applied to claim 27 above, and further in view of Michelson (U.S. Patent 4,968,298).

Froning, as applied to claim 27, discloses the claimed invention except for the step of introducing a flowable fill material into the interior region of the disc before introducing the lining. Michelson teaches to irrigate or wash out disc interspace after the

material from the disc has been removed, in order to remove any disc fragment and prevent inflammation of the neural elements and/or further surgery (see col. 1 line 5, through col. 2 line 46). It would have been obvious to one skill in the art at the time the invention was made to practice the method of Froning including the step of irrigate or wash out the interior region of the disc in view of Michelson, in order to be sure that no fragments of the nucleus pulposus are left inside the disc thus preventing inflammation and/or further surgery. With regard to claims 31-33, it would have been further obvious to one having ordinary skill in the art at the time the invention was made to irrigate or wash out the interior of the disc with naturally occurring extra-cellular matrix material, a slurry of at least one of saline, an antibiotic, a steroid, and a non-steroidal anti-inflammatory drug, or an autologous therapeutic agent, since it has been held to be within the general skill of a worker in the art to select a known material on the basis of its suitability for the intended use as a matter of obvious design choice. In re Leshin, 125 USPQ 416.

Conclusion

10. **THIS ACTION IS MADE FINAL.** Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the

shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Suba Ganesan whose telephone number is 571-272-3243. The examiner can normally be reached on M-F 8-5.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Corrine McDermott can be reached on 571-272-4754. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

BRIAN E. PELLEGRINO PRIMARY EXAMINER

SDG 5/21/2007